

**K183071 VLIFT-s Vertebral Body Replacement System**Jan 10, 2019  
66 days to decisionK183071 · Product code: **PLR** · Orthopedic  
Source: <https://www.510kdatabase.net/k183071/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Spinal Vertebral Body Replacement Device - Cervical (PLR)
Date received	Nov 5, 2018
Decision date	Jan 10, 2019
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Spine</b>
Location	Allendale, NJ, US
Contact	Renee Norby
510(k) history	74 submissions · 73 cleared · 2004-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k183071/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 16, 2026